DISPOSITION: On 3–30–60, pursuant to stipulation by the Government and by the claimant, Detergen Co., an order was entered directing the removal of the case to the Northern District of Oklahoma. Thereafter, interrogatories were served upon the claimant and subsequently answered.

On 5–5–61, the claimant having consented to a decree, judgment of condemnation was entered and the article was ordered released under bond to be brought into compliance with the law. On 1–15–62, the claimant having failed to post bond, an order for destruction of the article was entered.

6953. Red Rooster pills. (F.D.C. No. 46751. S. No. 42–883 T.)

QUANTITY: 26,400 tablets in a labeled bulk drum, 24,000 tablets in an unlabeled bulk drum, and 30 50-tablet btls., at Wyoming, Pa., in possession of Sanapac Co.

SHIPPED: 8–18–61, from Brooklyn, N.Y., by Manhattan Drug Co.

LABEL IN PART: (Drum) "S.F. 5073 Lot #4326 T Harmen Tablets Each tablet contains: * * * Ferrous Gluconate 100 mg. * * * Po. Ext. Passion Flower 100 mg. Po. Damiana 2 mg. Po. Nux Vomica 2 mg. * * * As a Hematinic and Bitter Tonic For Use in Iron Deficiency (Dietary) Anemias * * * Distributed by Manhattan Drug, Brooklyn 3, N.Y." and (btl.) "Sanapac's Red Rooster Pills 50 * * * Stimulant and Tonic A Dietary Supplement * * * Distributor The Sanapac Company, Wyoming, Penna. Formula * * * For Men and Women Only."

ACCOMPANYING LABELING: Window streamers reading in part "Red Rooster Pills give you that get up and go!"

RESULTS OF INVESTIGATION: The bottles described above were repacked by the dealer from bulk stock as described above.

LIBELED: 12–1–61, M. Dist. Pa.

CHARGE: 502(a)—when shipped and while held for sale, the labeling of the article contained false and misleading representations that the article was adequate and effective as a stimulant and tonic; to get up and go; and as a hematinic and bitter tonic for use in iron-deficiency anemia.

The article was alleged also to be misbranded under the provisions of the law applicable to foods, as reported in notices of judgment on foods.

DISPOSITION: 1–9–62. Default—destruction.

6954. Mineral bath. (F.D.C. No. 46604. S. No. 28–689 T.)

QUANTITY: 4 cases, 12 pkgs. each, at Kansas City, Mo.

SHIPPED: 4–13–61, from El Monte, Calif., by Oasis Co.

LABEL IN PART: (Pkg.) "New Hot Springs Mineral Way In Your Own Home Oasis Home Mineral Baths * * * Contain: Sodium Chloride, Sodium Sulphate, Aluminum Sulphate, Magnesium Sulphate, Sodium Tetraborate, Ammonium Aluminum Sulphate * * * Manufactured & Distributed by Oasis Co., El Monte, Calif."

ACCOMPANYING LABELING: Leaflets entitled "Oasis Home Mineral Baths" and display cards reading in part "Oasis Home Mineral Baths For External Use Only."

RESULTS OF INVESTIGATION: Examination showed the article to be in the form of crystals containing salts of sodium, aluminum, magnesium, and ammonia.

LIBELED: 10–26–61, W. Dist. Mo.

CHARGE: 502(a)—when shipped, the labeling contained false and misleading representations that the article would be effective, because of its mineral salt composition, as a treatment for relieving discomfort and pains of arthritis and rheumatism, relieving nervous tension, improving blood circulation, and that use of the article provides all the health benefits obtainable at health resorts.

DISPOSITION: 1–3–62. Default—destruction.

6955. Fountain-Facial with "Keroxylite." (F.D.C. No. 46768. S. No. 3–802 T.)

QUANTITY: 18,000 pkgs., 6 packets each, at Baltimore, Md.

SHIPPED: Between 4–14–61 and 5–5–61, from Wilmington, Del., by Packaging Services, Inc.

LABEL IN PART: (Pkg. and packet) "Six 9-gram Packets Fountain-Facial with 'Keroxylite' Skin Brightener, Cleanser, Antiseptic Each packet contains monopersulfate compound (Ozone*) sodium bicarbonate Alkyl-dimethyl-benzyl ammonium chloride * * * E. I. Du Pont De Nemours & Co., Inc. * * * Medically approved for cleansing and purifying the skin, and for treatment of acne, pimples, blackheads, inflamed hair follicles, and for inhibiting and removal of blemish-causing bacteria Directions * * * Hopkins Chemicals, Inc., Baltimore 2, Maryland."

ACCOMPANYING LABELING: Leaflet entitled "Fountain Facial with 'Keroxylite.'"

LIBELED: 12–5–61, Dist. Md.

CHARGE: 502(a)—when shipped, the labeling contained false and misleading representations that the article was an adequate and effective treatment for acne, pimples, blackheads, bacterial infections of the skin, and inflamed hair follicles.

DISPOSITION: 1–23–62. Default—destruction.

6956. Eterna 27 face cream. (F.D.C. No. 46629. S. Nos. 31–183/6 T.)

QUANTITY: 474 2-oz. jars at Riverside, Calif.; 435 2-oz. jars, 734 4-oz. jars, and 33 8-oz. jars, at San Bernardino, Calif.; and 419 2-oz. jars, 7 4-oz. jars, and 1 8-oz. jar, at Redlands, Calif., in possession of Sage's Complete Markets.

SHIPPED: Between 9–7–61 and 9–29–61, from Passaic, Metuchen, and Edison, N.J.

LABEL IN PART: (Jar) "'Eterna 27' Cream with Progenitin * * * Registered Trademark for Pregnenolone Acetate."

ACCOMPANYING LABELING: Newspaper advertisement in "San Bernardino Sun Telegram" and Riverside "Press Enterprise" both dated 10–15–61; leaflet "Eterna 27" reprint of an article in "Magazine of Wall Street and Business Analysts"; and counter display cards reading "This item [or "Product"] carries the Federal Pure Food and Drug Administration Seal of Approval Wash., D.C."

RESULTS OF INVESTIGATION: The accompanying labeling was prepared by the dealer and printed locally.

LIBELED: 11–14–61, S. Dist. Calif.

CHARGE: 502(a)—while held for sale, the labeling accompanying the article contained false and misleading representations that the article had been approved by the Food and Drug Administration.

DISPOSITION: 1–10–62. Consent—claimed by Sage's Complete Markets and relabeled.